We Claim:

- 1 1. (Original) A clear ibuprofen composition comprising:
- a. from about 15% to about 40% w/w of ibuprofen,
- b. from about 30% to about 70% w/w of polyethylene glycol,
- 4 c. from about 1% to about 10% w/w of a metal carbonate, and
- d. from about 1% to about 10% w/w of water.
- 1 2. (Cancelled)
- 1 3. (Original) The composition according to claim 1 wherein the polyethylene glycol
- 2 has an average molecular weight of about 300 to about 1000.
- 1 4. (Cancelled)
- 1 5. (Original) The composition according to claim 1 wherein the metal carbonate
- 2 comprises one or more of sodium bicarbonate, calcium carbonate, potassium
- 3 bicarbonate, sodium carbonate, potassium carbonate, magnesium carbonate,
- 4 magnesium bicarbonate, or mixtures thereof.
- 1 6. (Cancelled)
- 1 7. (Original) The composition according to claim 1 further comprising one or more
- 2 active ingredients, wherein the active ingredients comprise one or more of
- 3 glucosamine, pseudoephedrine, codeine, paracetamol, econazole, hydrocodone,
- 4 COX-2 inhibitors, alprazolam, dextromethorphan, chlorpheniramine, and
- 5 pharmaceutically acceptable salts thereof.
- 1 8. (Original) The composition according to claim 7 wherein the active ingredient
- 2 comprises pseudoephedrine and pharmaceutically acceptable salts thereof.
- 1 9. (Original) The composition according to claim 1 wherein the composition is filled
- 2 into soft gelatin capsules.
- 1 10. (Original) A process of preparing a clear ibuprofen composition, the process
- 2 comprising the steps of:
- a. dissolving one or more metal carbonates in water to form a solution,

b. adding ibuprofen and the solution of step (a) to polyethylene glycol with 4 5 optional heating, and 6 c. stirring to obtain a clear solution. 1 11. (Cancelled) 1 12. (Original) The process according to claim 10 wherein the polyethylene glycol has an average molecular weight of about 300 to about 1000. 2 (Cancelled) 1 13. (Original) The process according to claim 10 wherein the metal carbonate 1 14. comprises one or more of sodium bicarbonate, calcium carbonate, potassium 2 3 bicarbonate, sodium carbonate, potassium carbonate, magnesium carbonate, 4 magnesium bicarbonate, or mixtures thereof. 1 15. (Cancelled) (Original) The process according to claim 10 further comprising one or more 1 16. active ingredients, wherein the active ingredients comprise one or more of 2 3 glucosamine, pseudoephedrine, codeine, paracetamol, econazole, hydrocodone, 4 COX-2 inhibitors, alprazolam, dextromethorphan, chlorpheniramine, and 5 pharmaceutically acceptable salts thereof. 1 17. (Original) The process according to claim 16 wherein the active ingredient 2 comprises pseudoephedrine and pharmaceutically acceptable salts thereof. (Original) The process according to claim 10 further comprising filling the solution 1 18. 2 into a soft gelatin capsules. (Original) A soft gelatin capsule of ibuprofen, filled with a clear solution 1 19. 2 comprising: 3 a. from about 15% to about 40% w/w of ibuprofen, 4 b. from about 30% to about 70% w/w of polyethylene glycol, c. from about 1% to about 10% w/w of a metal carbonate, and 5 6 d. from about 1% to about 10% w/w of water. 1 20. (Original) The soft gelatin capsule of claim 19 wherein gelatin mass of the capsule 2 comprises gelatin, water, plasticizers, coloring agents and preservatives.

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(Previously Amended) The soft gelatin capsule of claim 20 wherein the plasticizers 1 21. 2 comprises sorbitol special solution and andrisorb. (Original) The soft gelatin capsule of claim 20 wherein the ratio of gelatin to water 1 22. varies from 1:0.75 to 1:0.92 and the ratio of gelatin to plasticizer varies from 2 3 1:0.35 to 1:0.48. (Original) The soft gelatin capsule according to claim 19 further comprising one or 1 23. 2 more active ingredients, selected from glucosamine, pseudoephedrine, codeine, paracetamol, econazole, hydrocodone, COX-2 inhibitors, alprazolam, 3 dextromethorphan, chlorpheniramine, and pharmaceutically acceptable salts 4 5 thereof. (Original) The soft gelatin capsule according to claim 23 wherein the one or more 1 24. 2 active ingredient is pseudoephedrine and pharmaceutically acceptable salts thereof. 25. (Cancelled) 1 1 26. (Cancelled) (Original) A clear ibuprofen-pseudoephedrine composition comprising: 1 27. a. from about 15% to about 40% w/w of ibuprofen, 1 b. from about 3% to about 6% w/w of pseudoephedrine or a pharmaceutically 2 3 acceptable salt thereof, c. from about 30% to about 70% w/w of polyethylene glycol, 4 5 d. from about 1% to about 10% w/w of a metal carbonate, and 6 e. from about 1% to about 10% w/w of water. 1 28. (Cancelled) 1 29. (Original) The composition according to claim 27 wherein the polyethylene glycol 2 has an average molecular weight of about 300 to about 1000. 30. (Cancelled) 1 (Original) The composition according to claim 27 wherein the metal carbonate 31. 1 2 comprises one or more of sodium bicarbonate, calcium carbonate, potassium

bicarbonate, sodium carbonate, potassium carbonate, magnesium carbonate,

magnesium bicarbonate, or mixtures thereof.

1	32.	(Cancelled)
1 2 3	33.	(Original) The composition according to claim 27 further comprising one or more active ingredients, wherein the active ingredient comprise one or more of glucosamine, codeine, paracetamol, econazole, hydrocodone, COX-2 inhibitors,
4 5		alprazolam, dextromethorphan, chlorpheniramine, and pharmaceutically acceptable salts thereof.
1 2	34.	(Original) The composition according to claim 27 wherein the composition is filled into soft gelatin capsules.
1 2	35.	(Original) A process of preparing a clear ibuprofen-pseudoephedrine composition comprising the steps of:
3		a. dissolving one or more metal carbonates in water to form a solution,
4 5		a. adding ibuprofen and the solution of step (a) to polyethylene glycol with optional heating,
6		b. stirring to obtain a clear solution, and
7 8		c. adding pseudoephedrine or a pharmaceutically acceptable salt thereof, and stirring to obtain a clear solution.
1 2	36.	(Original) The process according to claim 35 further comprising filling the solution of step (d) into a soft gelatin capsule.
1	37.	(Cancelled
1	38.	(Cancelled)
1 2	39.	(Original) A clear ibuprofen composition comprising: a. from about 15% to about 40% w/w of ibuprofen,
3		b. from about 30% to about 65% w/w of polyethylene glycol,
4		c. from about 1% to about 10% w/w of a metal carbonate,
5		d. from about 1% to about 15% w/w of a surfactant, and
6		e. from about 1% to about 10% w/w of water.
1	40.	(Cancelled)

- 1 41. (Original) The composition according to claim 39 wherein the polyethylene glycol 2 has an average molecular weight of about 300 to about 1000.
- 1 42. (Cancelled)
- 1 43. (Original) The composition according to claim 39 wherein the metal carbonate
- 2 comprises one or more of sodium bicarbonate, calcium carbonate, potassium
- 3 bicarbonate, sodium carbonate, potassium carbonate, magnesium carbonate,
- 4 magnesium bicarbonate, or mixtures thereof.
- 1 44. (Original) The composition according to claim 39 wherein the surfactant is a non-ionic hydrophilic surfactant.
- 1 45. (Original) The composition according to claim 44 wherein the non-ionic
- 2 hydrophilic surfactant comprises one or more of polyoxyethylene alkylethers,
- 3 polyethylene glycol fatty acids esters, polyethylene glycol glycerol fatty acid
- 4 esters, polyoxyethylene sorbitan fatty acid esters, polyoxyethylene-
- 5 polyoxypropylene block copolymers, polyglyceryl fatty acid esters,
- 6 polyoxyethylene glycerides, polyoxyethylene vegetable oils, and polyoxyethylene
- 7 hydrogenated vegetable oils.
- 1 46. (Original) The composition according to claim 39 further comprising one or more
- 2 active ingredients, wherein the active ingredients comprise one or more of
- glucosamine, pseudoephedrine, codeine, paracetamol, econazole, hydrocodone,
- 4 COX-2 inhibitors, alprazolam, dextromethorphan, chlorpheniramine, and
- 5 pharmaceutically acceptable salts thereof.
- 1 47. (Original) The composition according to claim 46 wherein the active ingredient
- 2 comprises pseudoephedrine and pharmaceutically acceptable salts thereof.
- 1 48. (Original) The composition according to claim 39 wherein the composition is filled
- 2 into soft gelatin capsules.
- 1 49. (Original) A process of preparing a clear ibuprofen composition, the process
- 2 comprising the steps of:
- a dissolving one or more metal carbonates in water to form a solution,
- b. preparing a solution of one or more surfactants in polyethylene glycol with
- 5 optional heating,
- 6 c. adding ibuprofen and the solution of step (a) to the solution of step (b), and

7 7 d. stirring to obtain a clear solution. 50. 1 (Cancelled) 1 51. (Original) The process according to claim 49 wherein the polyethylene glycol has 2 an average molecular weight of about 300 to about 1000. 1 52. (Cancelled) 1 53. (Original) The process according to claim 49 wherein the metal carbonate 2 comprises one or more of sodium bicarbonate, calcium carbonate, potassium 3 bicarbonate, sodium carbonate, potassium carbonate, magnesium carbonate, 4 magnesium bicarbonate, or mixtures thereof. 54. (Cancelled) 1 1 55. (Original) The process according to claim 49 wherein the surfactant comprises a 2 non-ionic hydrophilic surfactant. 1 56. (Original) The process according to claim 55 wherein the non-ionic hydrophilic 2 surfactant comprises one or more of polyoxyethylene alkylethers, polyethylene 3 glycol fatty acids esters, polyethylene glycol glycerol fatty acid esters, polyoxyethylene sorbitan fatty acid esters, polyoxyethylene-polyoxypropylene 4 5 block copolymers, polyglyceryl fatty acid esters, polyoxyethylene glycerides, 6 polyoxyethylene vegetable oils, and polyoxyethylene hydrogenated vegetable oils. 1 57. (Cancelled) 1 58. (Cancelled) 1 59. (Original) A clear ibuprofen-pseudoephedrine composition comprising: 2 from about 15% to about 40% w/w of ibuprofen, 3 b. from about 3% to about 6% w/w of pseudoephedrine, 4 c. from about 30% to about 65% w/w of polyethylene glycol, 5 d. from about 1% to about 10% w/w of a metal carbonate, 6 from about 1% to about 15% w/w of a surfactant, and 7 f. from about 1% to about 10% w/w of water.

1 60. (Cancelled)

(Original) The composition according to claim 59 wherein the polyethylene glycol 1 61. has an average molecular weight of about 300 to about 1000. 2 (Cancelled) The composition according to claim 61 wherein the polyethylene 62. 1 2 glycol has a molecular weight of about 400. (Original) The composition according to claim 59 wherein the metal carbonate 1 63. 2 comprises one or more of sodium bicarbonate, calcium carbonate, potassium 3 bicarbonate, sodium carbonate, potassium carbonate, magnesium carbonate, magnesium bicarbonate, or mixtures thereof. 4 1 64. (Cancelled) (Original) The composition according to claim 59 wherein the surfactant is a non-1 65. 2 ionic hydrophilic surfactant. (Original) The composition according to claim 65 wherein the non-ionic 1 66. 2 hydrophilic surfactant comprises one or more of polyoxyethylene alkylethers, polyethylene glycol fatty acids esters, polyethylene glycol glycerol fatty acid 3 esters, polyoxyethylene sorbitan fatty acid esters, polyoxyethylene-4 5 polyoxypropylene block copolymers, polyglyceryl fatty acid esters, polyoxyethylene glycerides, polyoxyethylene vegetable oils, and polyoxyethylene 6 hydrogenated vegetable oils. 7 (Original) The composition according to claim 59 further comprising one or more 1 67. 2 active ingredients, wherein the active ingredients comprise one or more of glucosamine, codeine, paracetamol, econazole, hydrocodone, COX-2 inhibitors, 3 alprazolam, dextromethorphan, chlorpheniramine, and pharmaceutically acceptable 4 5 salts thereof. (Original) The composition according to claim 59 wherein the composition is filled 68. 1 2 into soft gelatin capsules.

1 69. (Original) A process of preparing a clear ibuprofen-pseudoephedrine 2 composition comprising the steps of: 3 4 a. dissolving one or more metal carbonates in water to form a solution, 5 b. preparing a solution of one or more surfactants in polyethylene glycol with 6 optional heating, 7 c. adding ibuprofen and the solution of step (a) to the solution of step (b), 8 d. stirring to obtain a clear solution, and 9 e. adding pseudoephedrine or a pharmaceutically acceptable salt thereof to the solution of step (d) with continuous stirring to obtain a clear solution 10 1 70. (Cancelled) 1 71. (Original) The process according to claim 69 wherein the polyethylene glycol 2 has an average molecular weight of about 300 to about 1000. 72. (Original) The process according to claim 69 wherein the metal carbonate 1 2 comprises one or more of sodium bicarbonate, calcium carbonate, potassium 3 bicarbonate, sodium carbonate, potassium carbonate, magnesium carbonate, magnesium bicarbonate, or mixtures thereof. 4 1 .73. (Original) The process according to claim 69 wherein the surfactant comprises 2 a non-ionic hydrophilic surfactant. 74. (Original) The process according to claim 73 wherein the non-ionic hydrophilic 1 2 surfactant comprises one or more of polyoxyethylene alkylethers, polyethylene 3 glycol fatty acids esters, polyethylene glycol glycerol fatty acid esters, 4 polyoxyethylene sorbitan fatty acid esters, polyoxyethylene-polyoxypropylene block copolymers, polyglyceryl fatty acid esters, polyoxyethylene glycerides, 5 polyoxyethylene vegetable oils, and polyoxyethylene hydrogenated vegetable oils. 6 (Original) The process according to claim 69 further comprising one or more 1 75. 2 active ingredients, wherein the active ingredients comprise one or more of 3 glucosamine, codeine, paracetamol, econazole, hydrocodone, COX-2 inhibitors, alprazolam, dextromethorphan, chlorpheniramine, and pharmaceutically 4 5 acceptable salts thereof.

- 1 76. (Original) The process according to claim 69 further comprising filling the
- 2 solution of step (e) into a soft gelatin capsule.
- 1 77. (Cancelled)
- 1 78. (Cancelled)